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## Amendments to the claims:

Certain claims have been amended and others canceled below without disclaimer or prejudice to applicants' right to pursue the subject matter of these claims in a continuation application.

The following listing of claims will replace all prior versions, and listings, of claims in the application.

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## Listing of claims:

1. (Currently Amended) An improved process for the preparation of (S)-atenolol (1), comprising the steps of:

a) reacting a phenol of formula:

with an (R)-epichlorohydrin of formula:

in the presence of an alkali metal hydroxide and a quaternary ammonium salt as a phase transfer catalyst in an a solely aqueous solution at a temperature of -10°C to 0°C to obtain optically active intermediate glycidyl ether of formula:

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b) reacting the optically active intermediate glycidil ether (4) with isopropylamine at 10° to 40° C to obtain (S)-atenolol of formula

in high optical purity of >99% ee.

- 2. (Currently Amended) A The process as claimed in claim 1, wherein the alkali metal hydroxide is selected from sodium hydroxide or potassium hydroxide.
- 3. (Currently Amended) A <u>The</u> process as claimed in claim 1, wherein the amount of alkali metal hydroxide is 1 to 1.5 moles to 1 mole of the phenol (2).
- 4. (Currently Amended) A The process as claimed in claim 1, wherein the amount of (R)-epichlorohydrin is 1 to 3 moles to 1 mole of the phenol.
- 5. (Currently Amended) A The process as claimed in claim 1, wherein the quaternary ammonium salt has the formula R<sup>1</sup>R<sup>2</sup>R<sup>3</sup>R<sup>4</sup>N<sup>+</sup>X<sup>-</sup> wherein each of R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> are is the same or different and are is an alkyl group groups having 1 to

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16 carbon atoms selected from methyl, ethyl, propyl, butyl, phenyl or benzyl, X is a group selected from chlorine, bromine, iodine, hydrogen sulphate or hydroxyl.

- 6. (Currently Amended) A The process as claimed in claim 1, wherein the amount of quaternary ammonium salt is 0.001 to 2% by weight of phenol (2).
- 7. (Currently Amended) A The process as claimed in claim
  1 further comprising formation of chlorohydrin
  chlorohydrine (5) as a side product.
- 8. (Currently Amended) A The process as claimed in claim 7 ± further comprising reacting chlorohydrin chlorohydrine (5) with isolpropylamine isopropylamine at 10 to 40°C to obtain S-atenolol.
- 9. (New) The product of the improved process of claim 1, comprising (S)-atenolol of formula:

in high optical purity of >99% ee.